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IN THE CLAIMS

Please amend claim 67 as shown below. Please add new claims 68-81. The following listing of claims replaces all prior listings.

- 1-2. (Canceled).
- 3. (Previously presented) A targeted vesicle composition according to Claim 17 wherein:

$$X^{1}$$
 is $-C(=O)-NH-C(=O)-$;

$$X^{2}$$
 is $-C(=O)-$;

R¹ is acyl having from 16 to 20 carbons;

R³ is alkylene having from 1 to 3 carbons;

R⁴ is acyl having from 16 to 20 carbons;

R⁶ is a direct bond; and

R⁷ is lower alkylene.

- 4. (Previously presented) A targeted vesicle composition according to Claim 3 wherein:
 - R¹ is acyl having from 17 to 19 carbons;

R³ is methylene;

R⁴ is acyl having from 17 to 19 carbons; and

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R⁷ is ethylene.

5. (Canceled).

- 6. (Previously presented) A targeted vesicle composition according to Claim 17 wherein said hydrophilic polymer is selected from the group consisting of polyalkyleneoxides, polyvinyl alcohol, polyvinylpyrrolidones, polyacrylamides, polymethacrylamides, polyphosphazenes, poly(hydroxyalkylcarboxylic acids) and polyoxazolidines.
- 7. (Previously presented) A targeted vesicle composition according to Claim 6 wherein said hydrophilic polymer comprises a polyalkyleneoxide.
- 8. (Previously presented) A targeted vesicle composition according to Claim 7 wherein said hydrophilic polymer is selected from the group consisting of polyethylene glycol and polypropylene glycol.
- 9. (Previously presented) A targeted vesicle composition according to Claim 8 wherein said hydrophilic polymer is polyethylene glycol.
- 10. (Previously presented) A targeted vesicle composition according to Claim 8 wherein said hydrophilic polymer is PEG3400.
- 11. (Previously presented) A targeted vesicle composition according to Claim 17 wherein said targeting ligand comprises a peptide of the formula:

$$(Xaa)_n$$
— Yaa — Gly — Asp — $(Zaa)_m$

wherein:

each of m and n is independently an integer having value from 1 to 100;

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Xaa and Zaa are independently selected from the group consisting of natural amino acids and synthetic amino acids;

Yaa is selected from Arginine, Homoarginine, and Lysine-N-acetimidate; with the further proviso that when Xaa and Zaa are sulfur containing amino acids, Xaa and Zaa may be linked together via a disulfide linkage.

12. (Withdrawn) A targeted vesicle composition according to Claim 11, wherein:

Xaa is Glycine;

Yaa is Arginine;

Zaa is Serine;

n is 1, 2 or 3; and

m is 1.

13. (Withdrawn) A targeted vesicle composition according to Claim 12, wherein:

n is 3.

14. (Previously presented) A targeted vesicle composition according to Claim 11, wherein:

Xaa and Zaa comprise an amino acid independently selected from sulfur containing amino acids.

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wherein:

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15. (Previously presented) A targeted vesicle composition according to Claim 17 wherein said targeting ligand comprises a peptide of the following formula:

each x and y is independently an integer having value from 0 to 50;

each Saa is selected from the group consisting of natural and synthetic sulfur containing amino acids, wherein sulfur atoms in said sulfur containing amino acids are linked together by a disulfide bond, as represented by S—S;

each Xaa and Zaa are independently selected from the group consisting of natural amino acids and synthetic amino acids; and

Yaa is selected from Arginine, Homoarginine, and Lysine-N-acetimidate.

16. (Previously presented) A targeted vesicle composition according to Claim 15 wherein:

each Saa is independently selected from the group consisting of D-Cysteine, L-Cysteine, D-Penicillamine and L-Penicillamine.

17. (Previously presented) A targeted vesicle composition for therapeutic or diagnostic use *in vivo* comprising, in an aqueous carrier, gas filled liposomes comprising a phosphatidylcholine selected from the group consisting of dioleoylphosphatidylcholine, dimyristoylphosphatidylcholine, dipalmitoylphosphatidylcholine and

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distearoylphosphatidylcholine, wherein said liposomes further comprise a compound having the formula

$$R^{1} \xrightarrow{R^{2}} N \xrightarrow{N} R^{5}$$

$$R^{1} \xrightarrow{N} R^{3} - CH \xrightarrow{R^{6}} R^{6} - X^{1} - P \xrightarrow{R^{7}} R^{7} - X^{2} - T$$
(IV)

wherein:

each of X^1 and X^2 is independently a direct bond or a linking atom or group selected from the group consisting of $-C(=X^3)-$, $-C(=X^3)-$ N($R^8)-$ C($=X^3)-$ N($R^8)-$ C($=X^3)-$ N($=X^3)-$ N(

$$X^3$$
 is $-O-$ or $-S-$;

R¹ acyl having from 16 to 23 carbons;

R² is hydrogen or lower alkyl;

R³ is alkylene having from 1 to 10 carbons;

R⁴ acyl having from 16 to 23 carbons;

R⁵ is hydrogen or lower alkyl;

R⁶ is a direct bond:

R⁷ is a direct bond or alkylene having from 1 to 10 carbons;

R⁸ is hydrogen or lower alkyl;

P is a hydrophilic polymer; and

T is a targeting ligand which targets cells or receptors selected from the group consisting of myocardial cells, endothelial cells, epithelial cells, tumor cells and the glycoprotein GPIIbIIIa receptor.

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18-21. (Canceled).

- 22. (Previously presented) A targeted vesicle composition according to Claim 17 wherein said phosphatidylcholine comprises dipalmitoylphosphatidylcholine.
- 23. (Previously presented) A targeted vesicle composition according to Claim 17 further comprising a phosphatidylethanolamine selected from the group consisting of dipalmitoyl-phosphatidylethanolamine, dioleoylphosphatidylethanolamine, N-succinyldioleoyl-phosphatidylethanolamine and 1-hexadecyl-2-palmitoylglycerophosphoethanolamine.
- 24. (Original) A targeted vesicle composition according to Claim 23 wherein said phosphatidylethanolamine comprises dipalmitoylphosphatidylethanolamine.
- 25. (Previously presented) A targeted vesicle composition according to Claim 17 further comprising dipalmitoylphosphatidic acid.
- 26. (Original) A targeted vesicle composition according to Claim 17, wherein said vesicles comprise a gas selected from the group consisting of perfluorocarbons and sulfur hexafluoride.
- 27. (Original) A targeted vesicle composition according to Claim 26 wherein said perfluorocarbon gas is selected from the group consisting of perfluoromethane, perfluoroethane, perfluoropropane, perfluorobutane and perfluorocyclobutane.
- 28. (Original) A targeted vesicle composition according to Claim 27 wherein said perfluorocarbon gas is selected from the group consisting of perfluoropropane and perfluorobutane.

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29. (Original) A targeted vesicle composition according to Claim 28 wherein said perfluorocarbon gas comprises perfluorobutane.

- 30. (Original) A targeted vesicle composition according to Claim 17 wherein said gas is derived, at least in part, from a gaseous precursor.
- 31. (Original) A targeted vesicle composition according to Claim 30 wherein said gaseous precursor has a boiling point of greater than about 37°C.
- 32. (Original) A targeted vesicle composition according to Claim 31 wherein said gaseous precursor comprises a perfluorocarbon.
- 33. (Original) A targeted vesicle composition according to Claim 32 wherein said perfluorocarbon is selected from the group consisting of perfluoropentane and perfluorohexane.
- 34. (Original) A targeted vesicle composition according to Claim 17 wherein said vesicles further comprise a bioactive agent that is different from said gas and said compound.
- 35. (Original) A targeted vesicle composition according to Claim 34 wherein said bioactive agent comprises a therapeutic agent selected from the group consisting of genetic material, dihydroergotamine, heparin sulfate, tissue plasminogen activator, streptokinase, urokinase, hirudin, and mixtures thereof.

36-60. (Canceled).

61. (Previously presented) A targeted vesicle composition according to Claim 4 wherein:

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each of R¹ and R⁴ is acyl of 18 carbons.

62. (Canceled)

63. (Previously presented) A targeted vesicle composition according to Claim 4 wherein:

R¹ is an acyl of 18 carbons.

- 64. (Previously presented) A targeted vesicle composition according to Claim 17, wherein said targeting ligand T is a peptide having from 3 to 20 amino acids.
- 65. (Previously presented) A targeted vesicle composition according to Claim 64, wherein said peptide is cyclized by a linkage selected from the group consisting of sidechain to-sidechain covalent linkages, end-to-sidechain covalent linkages, and end-to-end covalent linkages.
- 66. (Previously presented) A targeted vesicle composition for therapeutic or diagnostic use *in vivo* comprising, in an aqueous carrier, gas filled liposomes comprising a phosphatidylcholine selected from the group consisting of dioleoylphosphatidylcholine, dimyristoylphosphatidylcholine, dipalmitoylphosphatidylcholine and distearoylphosphatidylcholine, wherein said liposomes further comprise a compound having the formula

$$R^{2}$$
 $N \longrightarrow R^{5}$
 $R^{1} \longrightarrow N \longrightarrow R^{3} \longrightarrow CH \longrightarrow R^{6} \longrightarrow X^{1} \longrightarrow P \longrightarrow R^{7} \longrightarrow X^{2} \longrightarrow T$

wherein:

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$$X^1$$
 is $-C(=X^3)-N(R^8)-$;

$$X^{2}$$
 is $C(=X^{3})$;

$$X^3$$
 is O;

each of R1 and R4 is acyl having 18 carbons;

each of R², R⁵ and R⁸ is H;

each of R³ and R⁷ is ethylene;

R⁶ is a direct bond;

P is PEG-3400; and

T comprises a peptide having the sequence CRGDC, wherein the two cysteines are linked together via a disulfide linkage.

- 67. (Currently amended) The A targeted vesicle composition according to Claim 66 81, further comprising wherein said bioactive agent is urokinase.
- 68. (New) The targeted vesicle composition according to Claim 66, wherein said phosphatidylcholine comprises dipalmitoylphosphatidylcholine.
- 69. (New) The targeted vesicle composition according to Claim 66, further comprising a phosphatidylethanolamine selected from the group consisting of dipalmitoyl-phosphatidylethanolamine, dioleoylphosphatidylethanolamine, N-succinyldioleoyl-phosphatidylethanolamine and 1-hexadecyl-2-palmitoylglycerophosphoethanolamine.

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70. (New) The targeted vesicle composition according to Claim 69, wherein said phosphatidylethanolamine comprises dipalmitoylphosphatidylethanolamine.

- 71. (New) The targeted vesicle composition according to Claim 66, further comprising dipalmitoylphosphatidic acid.
- 72. (New) The targeted vesicle composition according to Claim 66, wherein said vesicles comprise a gas selected from the group consisting of perfluorocarbons and sulfur hexafluoride.
- 73. (New) The targeted vesicle composition according to Claim 72, wherein said perfluorocarbon gas is selected from the group consisting of perfluoromethane, perfluoropropane, perfluorobutane and perfluorocyclobutane.
- 74. (New) The targeted vesicle composition according to Claim 73, wherein said perfluorocarbon gas is selected from the group consisting of perfluoropropane and perfluorobutane.
- 75. (New) The targeted vesicle composition according to Claim 74, wherein said perfluorocarbon gas comprises perfluorobutane.
- 76. (New) The targeted vesicle composition according to Claim 66, wherein said gas is derived, at least in part, from a gaseous precursor.
- 77. (New) The targeted vesicle composition according to Claim 76, wherein said gaseous precursor has a boiling point of greater than about 37°C.
- 78. (New) The targeted vesicle composition according to Claim 76, wherein said gaseous precursor comprises a perfluorocarbon.

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The targeted vesicle composition according to Claim 78, wherein 79. (New) said perfluorocarbon is selected from the group consisting of perfluoropentane and perfluorohexane.

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- The targeted vesicle composition according to Claim 66, wherein 80. (New) said vesicles further comprise a bioactive agent that is different from said gas and said compound.
- The targeted vesicle composition according to Claim 80, wherein 81. (New) said bioactive agent comprises a therapeutic agent selected from the group consisting of genetic material, dihydroergotamine, heparin sulfate, tissue plasminogen activator, streptokinase, urokinase, hirudin, and mixtures thereof.